

REPORT DOCUMENTATION PAGE

Form Approved
OMB No. 0704-0188

Public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to Washington Headquarters Services, Directorate for Information Operations and Reports, 1215 Jefferson Davis Highway, Suite 1204, Arlington, VA 22202-4302, and to the Office of Management and Budget, Paperwork Reduction Project (0704-0188), Washington, DC 20503.

PLEASE DO NOT RETURN YOUR FORM TO THE ABOVE ADDRESS.

1. REPORT DATE (DD-MM-YYYY) 01-11-2004			2. REPORT TYPE Final Report		3. DATES COVERED Sept 1999 - December 2001	
4. TITLE AND SUBTITLE Instant Cell Analysis, BioSPICE, Cellular Devices and Exquisite Detection: Towards an Interactive Biology					5a. CONTRACT NUMBER	
					5b. GRANT NUMBER N00014-99-F-0458	
					5c. PROGRAM ELEMENT NUMBER	
6. AUTHOR(S) Arkin,Adam, P					5d. PROJECT NUMBER	
					5e. TASK NUMBER	
					5f. WORK UNIT NUMBER	
7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES) University of California, Berkeley mailcode 5230 Calvin Lab Berkeley, CA 94720-5230					8. PERFORMING ORGANIZATION REPORT NUMBER	
9. SPONSORING/MONITORING AGENCY NAME(S) AND ADDRESS(ES) Office of Naval Research 800 N. Quincy Street Arlington, VA 22217-5000					10. SPONSOR/MONITOR'S ACRONYM(S) ONR	
					11. SPONSOR/MONITOR'S REPORT NUMBER(S)	
12. DISTRIBUTION/AILABILITY STATEMENT Distribution Unlimited						
13. SUPPLEMENTARY NOTES						
<div style="text-align: right; font-size: 2em; font-weight: bold;">20041112 024</div>						
14. ABSTRACT The overall objective of this project was to rapidly find molecular reagents to interfere or bind to every gene product in the <i>Saccharomyces cerevisiae</i> through the development of new screening and data analytical technologies. Two experimental approaches were taken. Ron Davis developed the haploinsufficiency profiling (HIP) technology wherein two libraries of yeast were constructed one containing strains missing one copy the other strains missing both copies of every gene in yeast (that could be knock out). In place of its missing gene, each strain was engineered with a DNA bar code to identify it. Affymetrix chips were used to measure the relative populations of each strain under different chemical perturbations. Roger Brent worked on build protein aptamers to bind gene products. Arkin's role was in the development of analyses for the Haploinsufficiency data.						
15. SUBJECT TERMS haploinsufficiency, yeast, affymetrix chips						
16. SECURITY CLASSIFICATION OF:			17. LIMITATION OF ABSTRACT		18. NUMBER OF PAGES	
a. REPORT	b. ABSTRACT	c. THIS PAGE				
unclass	unclass	unclass	UL	self expl	19a. NAME OF RESPONSIBLE PERSON Adam P. Arkin	
					19b. TELEPHONE NUMBER (Include area code) 510-495-2366	

Grant #: N00014-99-F-0458

Principal Investigator: Adam P. Arkin

Institution: E.O. Lawrence Berkeley National Laboratory

Grant Title: Instant Cell Analysis & BioSPICE

Award Period: 1 September 1999 - 31 December 2001

Objective: The overall objective of this project was to rapidly find molecular reagents to interfere or bind to every gene product in the *Saccharomyces cerevisiae* through the development of new screening and data analytical technologies.

Approach: Two experimental approaches were taken. Ron Davis developed the haploinsufficiency profiling (HIP) technology wherein two libraries of yeast were constructed one containing strains missing one copy the other strains missing both copies of every gene in yeast (that could be knock out). In place of its missing gene, each strain was engineered with a DNA bar code to identify it. Affymetrix chips were used to measure the relative populations of each strain under different chemical perturbations. Roger Brent worked on build protein aptamers to bind gene products. Arkin's role was in the development of analyses for the Haploinsufficiency data.

Accomplishments: Our role on the project was to aid in the quality analysis and quality control of the HIP pipeline and to develop the databases and analyses for this data. During the course of this project we developed the database and the initial analyses that led to proof of the quality and reproducibility of the method at moderate scale screening[1]. We discovered a number of interesting facts: The gene necessary for surviving a perturbation to the environment were not represented by the set of genes that changed gene expression (adding a caveat to the power gene expression microarray techniques) and that we are able to determine accurately the molecular target and gross mechanism of action of each of the drugs or environmental exposures the cell is subjected to. The success of the project during this grant was parlayed into a larger NIH grant in which we have screened hundreds of small molecules for the effects on the population--the database is now large, the analyses of the data are very sophisticated leading to very robust prediction of molecular targets and we have developed new algorithms for classification of genes and chemical perturbants that allow deduction of

orthogonal perturbation sets and prediction of the effects of new chemicals[2, 3].

Conclusions: HIP profiling technology is an effective method for rapidly screening large chemical libraries to infer the action of each molecule on the proteins of yeast and the attendant mechanisms of action. We were able not only to correctly identify the known target of a number of antifungals but discover the mechanisms of less characterized members of the library and discover the pleiotropic effects that make some drugs less specific than others.

Significance: Apart from developing and hardening a entirely new technology for screening the activity of new molecules and identifying their protein targets--this method is now being adapted to screen for new classes of druggable proteins for use in human therapeutics and for deducing networks of interaction in the yeast cell.

Patent Information: No patents have been filed related to this research

Award Information: TR100, Time Magazine Top Innovator

Publication and Abstracts:

1. Giaever, G., et al., *Functional profiling of the Saccharomyces cerevisiae genome*. Nature, 2002. 418(6896): p. 387-91.
2. Giaever, G., et al., *Chemogenomic profiling: identifying the functional interactions of small molecules in yeast*. Proc Natl Acad Sci U S A, 2004. 101(3): p. 793-8.
3. Flaherty, P., et al., *A Latent Variable Model for Chemogenomic Profiling*. Bioinformatics, 2004. Accepted Conditionally.

Numerous talks were given during the period of which this topic was a portion. A full list of invited talks during this project period is available upon request.